AMENDMENTS TO THE CLAIMS

1. **(Currently amended)** A flavivirus having a phenotype in which the viral genome is modified by the introduction of a mutation, singly or in combination, taken from the group consisting of the mutations of ~~any of Table 1-37, preferably~~ Table 37, wherein said mutation is charge-cluster-to-alanine mutations 200, 201.

2. **(Original)** The flavivirus of claim 1, further comprising the Δ30 mutation.

3. **(Original)** The flavivirus of claim 1, wherein the flavivirus is a dengue virus type 1.

4. **(Original)** The flavivirus of claim 1, wherein the flavivirus is a dengue virus type 2.

5. **(Original)** The flavivirus of claim 1, wherein the flavivirus is a dengue virus type 3.

6. **(Original)** The flavivirus of claim 1, wherein the flavivirus is a dengue virus type 4.

7. **(Original)** The flavivirus of claim 1, wherein the flavivirus is a chimeric virus.

8. **(Original)** The chimeric virus of claim 7 having a dengue 1 backbone.

9. **(Original)** The chimeric virus of claim 7 having a dengue 2 backbone.

10. **(Original)** The chimeric virus of claim 7 having a dengue 3 backbone.

11. **(Original)** The chimeric virus of claim 7 having a dengue 4 backbone.

12. **(Withdrawn)** The flavivirus of claim 1, wherein the phenotype is temperature sensitivity in Vero cells or the human liver cell line HuH-7.

13. **(Withdrawn)** The flavivirus of claim 1, wherein the phenotype is host-cell restriction in mosquito cells or the human liver cell line HuH-7.

14. **(Withdrawn)** The flavivirus of claim 1, wherein the phenotype is host-cell adaptation for improved replication in Vero cells.

15. **(Withdrawn)** The flavivirus of claim 1, wherein the phenotype is attenuation in mice.

16. **(Withdrawn)** A pharmaceutical composition comprising a pharmacologically acceptable vehicle and a flavivirus according to any of claims 1-15.

17. **(Withdrawn)** A kit comprising a pharmaceutical composition according to claim 16 in a pack or dispenser device and instructions for administration.

18. **(Withdrawn)** A method of producing neutralizing antibodies against dengue virus comprising the administration of a therapeutically effective amount of a pharmaceutical composition comprising a pharmacologically acceptable vehicle and a flavivirus according to any of claims 1-15.

19. **(Withdrawn)** The method of claim 18, wherein administration is by subcutaneous, intradermal, or intramuscular injection.

20. **(Withdrawn)** A tetravalent vaccine comprising a pharmacologically acceptable vehicle and a flavivirus according to any of claims 1-15.

21. **(Withdrawn)** An live attenuated vaccine comprising a pharmacologically acceptable vehicle and a flavivirus according to any of claims 1-15.

22. **(Withdrawn)** The live attenuated vaccine of claim 21 in unit dosage form having from about 10^2 –10^9 plaque forming units (PFU)/ml.

23. **(Withdrawn)** An inactivated vaccine comprising a pharmacologically acceptable vehicle and a flavivirus according to any of claims 1-15.

24. **(Withdrawn)** The inactivated vaccine of claim 23 in unit dosage form having from about 0.1 to 50 μg of E protein/ml.

25. **(Withdrawn)** A cDNA molecule encoding a flavivirus according to any of claims 1-15.

26. **(Withdrawn)** An RNA molecule encoding a flavivirus according to any of claims 1-15.

27. **(Withdrawn)** A method of preparing a flavivirus comprising (a) synthesizing full-length viral genomic RNA in vitro using a cDNA molecule that encodes a flavivirus according to any of claims 1-15; (b) transfecting cultured cells with the viral genomic RNA to produce virus; and (c) isolating the virus from the cultured cells.

28. **(Withdrawn)** A method of making a pharmaceutical composition comprising combining a pharmacologically acceptable vehicle and a flavivirus according to any of claims 1-15.

29. **(Withdrawn)** A method of identifying a mutation that restricts replication in human liver cells comprising (a) introducing mutations into a dengue virus genome to make mutant viruses; (b) screening the mutant viruses for a phenotype characterized by host-cell restriction in human liver cells; and (c) determining the genetic basis for the phenotype by direct sequence analysis of the virus genome.

30. **(Withdrawn)** A method of identifying a mutation that promotes growth in Vero cells comprising (a) introducing mutations into a dengue virus genome to make mutant viruses; (b) screening the mutant viruses for a phenotype characterized by host-cell adaptation for improved replication in Vero cells; and (c) determining the genetic basis for the phenotype by direct sequence analysis of the virus genome.

31. **(Withdrawn)** A method of assembling a menu of mutations for use in fine-tuning the attenuation and growth characteristics of recombinant dengue viruses comprising (a) introducing mutations into a dengue virus genome to make mutant viruses; (b) screening the mutant viruses for a phenotype characterized by temperature sensitivity in Vero cells or human liver cells, host cell restriction in mosquito cells or human liver cells, host-cell adaptation for improved replication in Vero cells, or attenuation in mice; (c) determining the genetic basis for the phenotype by direct sequence analysis of the virus genome; and (d) performing multiple iterations of steps (a) – (c), whereby a menu of mutations is assembled.

32. **(Withdrawn)** The method of any of claims 29-30 further comprising introducing the mutation identified by said method into a recombinant dengue virus, and characterizing the resulting phenotype.